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RAW SEQUENCE LISTING
PATENT APPLICATION: US/09/484,577A

DATE: 02/13/2001
TIME: 14:31:57

Input Set : A:\029001.txt
Output Set: N:\CRF3\02132001\I484577A.raw

4 <110> APPLICANT: Gordon, Lynn K.
5 Goodlick, Lee
6 Goldman, Melissa
8 <120> TITLE OF INVENTION: NOVEL GENES AND POLYPEPTIDES FOR THE
9 DIAGNOSIS OF GIANT CELL ARTERITIS
12 <130> FILE REFERENCE: 07419-029001
14 <140> CURRENT APPLICATION NUMBER: 09/484,577A
15 <141> CURRENT FILING DATE: 2000-01-18
17 <160> NUMBER OF SEQ ID NOS: 98
19 <170> SOFTWARE: FastSEQ for Windows Version 4.0
21 <210> SEQ ID NO: 1
22 <211> LENGTH: 682
23 <212> TYPE: DNA
24 <213> ORGANISM: Homosapiens
26 <220> FEATURE:
27 <223> OTHER INFORMATION: artificially generated nucleic acid
29 <400> SEQUENCE: 1
30 gatcccccgct ttcgcgggta tgacagcggt actcaattca cgcgcccgca tggcagcgaa 60
31 ctaaacggag gatctcacga acatccgctc caaccccgac accacgtcc cccgcgtcac 120
32 gacaggctcg ctggccctct cgcgcaagtt ctttgcataat cctgaggccg cgcgcacat 180
33 cgcgttccc ttgcgcgaga tcatcctgtc cgaggccgc ggcgagccga acctgcccgt 240
34 ctatgacacc tcggggccct acaccgatcc ggcgtgcacg atcgacgtca acaggccct 300
35 gccgcgcaat cgccctcgct gggtaaaggaa acggggccgc gtcgaggaat atcaggccgc 360
36 accatcaagc cggaggacaa cggcaatgtc ggcgcattccc acggccgcca ggcgttacc 420
37 ggaccacaaa gcccgtgcgc ggctcgacgg cacaagatca cccactcgag ttcgcgcgc 480
38 cggcattata ccaaggagat gatctacgtc gccgagcgtg agaatctgg cgcaagcagc 540
39 agctgagcgc gccgaggccg gctgcgcacg gaagagttt ggcgcgcgg tgccggctta 600
40 ttacgcccggaa atttgcgca agagatcgcg cggccgcatt tatttcctttaaaaattaaaca 660
41 ttggcgagct tgaaccgatg aa 682
43 <210> SEQ ID NO: 2
44 <211> LENGTH: 92
45 <212> TYPE: PRT
46 <213> ORGANISM: Homosapiens
48 <400> SEQUENCE: 2
49 Leu Pro Ala Val Thr Thr Gly Ser Leu Pro Ser Ser Arg Lys Phe Phe
50 1 5 10 15
51 Ala Ile Pro Glu Ala Ala Pro Asp Ile Arg Val Pro Leu Arg Glu Ile
52 20 25 30
53 Ile Leu Ser Glu Gly Ala Gly Glu Pro Asn Leu Pro Val Tyr Asp Thr
54 35 40 45
55 Ser Gly Pro Tyr Thr Asp Pro Ala Val Thr Ile Asp Val Asn Ser Gly
56 50 55 60
57 Leu Pro Arg Asn Arg Leu Ala Trp Val Lys Glu Arg Gly Gly Val Glu
58 65 70 75 80
59 Glu Tyr Gln Ala Ala Pro Ser Ser Arg Arg Thr Thr
60 85 90
62 <210> SEQ ID NO: 3

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63 <211> LENGTH: 501
64 <212> TYPE: DNA
65 <213> ORGANISM: Homosapiens
67 <220> FEATURE:
68 <223> OTHER INFORMATION: Synthetically generated nucleic acid
70 <400> SEQUENCE: 3
71 actctccagc ctctcaccga ggtatgaagtc ggctcgtaa gttgggtcgaa tcgggggcaa 60
72 aacccgggac gagctggct tcctgcggc cgcgcctcgaa attgtcgaga cgccgcac 120
73 tcccaccgag agactcacgg cgcgccttgct tgctgccttg ttctactgcg cgcgtggcgtg 180
74 ggcgggtctc ggcaggatcg acatcggtgc ttctgcaccc agaaagatcg tgccggcgaa 240
75 cgcgttaaag ctgggttcagc cgctcgaggc cggcgtgggt cggccactc atgtccgcga 300
76 tggccaaacc gtcaaggccg gcgagattct gatcgagtcg gatccatcg cgggtgggt 360
77 ggtatgttcg ccccgatcaga ggtccatcac ggtgtccggc ccccacggat cgcacaccca 420
78 tcttgcac ctttcttcac cgacgagtca cccggcagtt gccgatattg cgtgatctta 480
79 tcagaatgcg gcgatgtatca t 501
81 <211> SEQ ID NO: 4
82 <211> LENGTH: 124
83 <212> TYPE: PRT
84 <213> ORGANISM: Homosapiens
86 <400> SEQUENCE: 4
87 Leu Ser Ser Leu Ser Pro Arg Met Lys Ser Ala Arg Glu Val Val Ala
88 1 5 10 15
89 Val Gly Gly Lys Thr Arg Asp Glu Leu Ala Phe Leu Pro Ala Ala Leu
90 20 25 30
91 Glu Ile Val Glu Thr Pro Pro Ser Pro Thr Ala Arg Leu Thr Ala Ala
92 35 40 45
93 Leu Leu Ala Ala Leu Phe Tyr Cys Ala Val Ala Trp Ala Gly Leu Gly
94 50 55 60
95 Arg Ile Asp Ile Val Ala Ser Ala Ser Arg Lys Ile Val Pro Gly Asp
96 65 70 75 80
97 Arg Val Lys Leu Val Gln Pro Leu Glu Val Gly Val Val Arg Ala Thr
98 85 90 95
99 His Val Arg Asp Gly Gln Thr Val Lys Ala Gly Glu Ile Leu Ile Glu
100 100 105 110
101 Leu Asp Pro Phe Ala Gly Val Asp Val Ala Thr
102 115 120
104 <210> SEQ ID NO: 5
105 <211> LENGTH: 747
106 <212> TYPE: DNA
107 <213> ORGANISM: Homosapiens
109 <220> FEATURE:
110 <223> OTHER INFORMATION: Artificially generated nucleic acid
112 <400> SEQUENCE: 5
113 accgacgtcg actatccatg aacggatccc tgcaacgaca tcgtcgtaa ggcctatgaa 60
114 ggcgtcgccg ccgtgctcg tggcacgcag tcgctccaca ccaactcgat cgacgaggcg 120
115 atcgcgctgc cgattgactt ctccggccgg atcgccgcac acaccagctg atccagcgc 180
116 acggagacaga cgtcacggac gcggtcgaca ctctggcgaa gtcctactac gtggagcgcc 240
117 tgacggatga cctcgccaaag cggggctggg agctgatgaa agaggtcgag aagatgggtg 300
118 gcatggcgca ggcgatcgcc accgggttggc cgaaggccct gatcgagcaa tctcgacgc 360

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119 aaaagcaggc cgcgatcgat cgccgcgatc aggtgatcgt gggcgtgaac cgctaccggc 420
 120 ccgaacacgga gcaaccgatc gacattattg agatcgacaa ctgcacggtt cgggcctccc 480
 121 agatccgggt tctcgccgaa atcgaaaagg cgccgtgatc aaggaagggtt gagtccgcgc 540
 122 tcggggagct ggcgtgtatt gcccgcacgg gtgagggaaa tctgctggct gcagcgaccg 600
 123 agcccgctcg cgccgcggct accgtcgaaa agatgtccga cgccatgcgg caagcattcg 660
 124 gcgaccacga ggcgggtcccg gaggtgtgtt cggacgttta cggccgtgcc tatggcacgg 720
 125 atccgttcat ggatagtcga cgtcggt 747
 127 <210> SEQ ID NO: 6
 128 <211> LENGTH: 48
 129 <212> TYPE: PRT
 130 <213> ORGANISM: Homosapiens
 132 <400> SEQUENCE: 6
 133 Asp Pro Cys Asn Asp Ile Val Arg Thr Ala Tyr Glu Ala Leu Ala Ala
 134 1 5 10 15
 135 Val Leu Gly Gly Thr Gln Ser Leu His Thr Asn Ser Phe Asp Glu Ala
 136 20 25 30
 137 Ile Ala Leu Pro Ile Asp Phe Ser Ala Arg Ile Ala Arg Asn Thr Ser
 138 35 40 45
 140 <210> SEQ ID NO: 7
 141 <211> LENGTH: 301
 142 <212> TYPE: DNA
 143 <213> ORGANISM: Homosapiens
 145 <220> FEATURE:
 146 <223> OTHER INFORMATION: Synthetically generated nucleic acid
 148 <400> SEQUENCE: 7
 149 actctccagc ctctcacgga ggatcatcgaa cgacattaaag cagctggccg acaacggcgt 60
 150 ggcgcgaattc acgctgatcg gacagaatgt caacgcctac cacggccggag ggcccgacgg 120
 151 ccgcgtctgg ccgcctcgca aattgctgca gcgactcgcg gacattccag gcgtcatcg 180
 152 gctgcgttat tcgatcagcc atccgcgcga cgtcgacgac agcctgatcg cccgcgcacg 240
 153 cgatttgccc ggactgtatgc cgttcgtgca cctgcccgtt caatcgaaaa cggaccggat 300
 154 c 301
 156 <210> SEQ ID NO: 8
 157 <211> LENGTH: 91
 158 <212> TYPE: PRT
 159 <213> ORGANISM: Homosapiens
 161 <400> SEQUENCE: 8
 162 Ile Ile Asp Asp Ile Lys Gln Leu Ala Asp Asn Gly Val Arg Glu Phe
 163 1 5 10 15
 164 Thr Leu Ile Gly Gln Asn Val Asn Ala Tyr His Gly Gly Gly Pro Asp
 165 20 25 30
 166 Gly Arg Val Trp Pro Leu Gly Lys Leu Leu Gln Arg Leu Ala Asp Ile
 167 35 40 45
 168 Pro Gly Val Met Arg Leu Arg Tyr Ser Ile Ser His Pro Arg Asp Val
 169 50 55 60
 170 Asp Asp Ser Leu Ile Ala Ala His Arg Asp Leu Pro Gly Leu Met Pro
 171 65 70 75 80
 172 Phe Val His Leu Pro Val Gln Ser Gly Ala Asp
 173 85 90
 175 <210> SEQ ID NO: 9

RAW SEQUENCE LISTING

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Input Set : A:\029001.txt

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176 <211> LENGTH: 620
 177 <212> TYPE: DNA
 178 <213> ORGANISM: Homosapiens
 180 <220> FEATURE:
 181 <221> NAME/KEY: misc_feature
 182 <222> LOCATION: (0)...(0)
 183 <223> OTHER INFORMATION: N=A,T,C or G
 185 <400> SEQUENCE: 9

W--> 186	actctccanc	ctctcaccgga	ggatcagaat	aggtaaagag	cgaagacacc	gagaacgtct	60
187	ggccttgaac	ggacacgctg	cttgagttgg	tcggggtcac	caccgaccc	gtgtccaccg	120
W--> 188	gcgcaagtac	ngtggaaagca	cttgaccatg	atcccagacg	gtggcgat	ccgcgcggac	180
W--> 189	ccacancgtt	cccgccccc	accggattga	tagctcagcg	acaccagctg	ggctgcccgt	240
W--> 190	acgttattgt	gctgggttng	tgcaagtgc	accccgctca	agacaaantg	gccgcacactg	300
W--> 191	tgcggctgtc	ccaaacgtca	tattgggtcg	cagcaactgtc	gaacggatca	ctgtangtgc	360
W--> 192	acagcgacna	anccgcata	ctctngccgt	ggggcgcaac	gatgtttnac	accgtctcaa	420
W--> 193	cgggtaccgt	gtnagggga	ncatttacng	ggaaagcatt	cgaccactcc	cccacaccgt	480
W--> 194	gccccgcattt	gcccgcattt	ctttcattga	tatgtccacg	tcggtnngnc	tttaagcngg	540
W--> 195	cggcaaccgc	ggtgnagctn	cacttttgt	tccttttatt	ganggttaat	ttgcgcgtt	600
W--> 196	tggncgtaan	tntttngaan					620
198 <210>	SEQ ID NO:	10					
199 <211>	LENGTH:	662					
200 <212>	TYPE:	DNA					
201 <213>	ORGANISM:	Homosapiens					
203 <220>	FEATURE:						
204 <221>	NAME/KEY:	misc_feature					
205 <222>	LOCATION:	(0)...(0)					
206 <223>	OTHER INFORMATION:	N=A,T,C or G					
208 <400>	SEQUENCE:	10					
209	gatccgacca	gcaatcaggc	ggagctgcag	cacctgaaaa	acgacccttct	ctccgcactg	60
210	ctgggtattt	cacgcaaccg	ctctgcgtt	ggcgaaaaac	accgacgcgc	ttgaaggctt	120
211	accggacgac	acggcccaag	ccttgattcg	aatgcacatcg	gagtaacttgc	gcagtcagga	180
212	ttccgagcag	cgcgcac	tgtccgaact	ggatcagcaa	cgggtgcaga	aggtcgccga	240
W--> 213	gaccaggacg	atcgacgcca	gcacgcgaa	gattgaagct	ttgctgcgt	gctgcaggan	300
214	cgggtcgaaa	ttcgcaagta	cctggccgac	agggagatcg	gctcaaaagct	gcaatattcg	360
215	caggaactcc	aggaacttgg	cgggatgcag	caggacatcc	tggtgcacg	gagcaaaagct	420
W--> 216	cggggaaaacc	aatgcggntt	gtcgccgcac	ttcgacgaaa	acccgcggna	agttcgtct	480
W--> 217	nnngaataacc	ggcacccgnc	tgttccnacg	atcttggccc	aaggggacgc	aaaaaaggc	540
W--> 218	cggcaagnc	tcaaaggacc	aagggnngttt	taaaanccga	gcacccggga	cccaaccttt	600
W--> 219	aaaaancntt	ggcgcccccc	attcgacggn	gtggnggca	caaattggc	cgngccccat	660
220	tt						662
222 <210>	SEQ ID NO:	11					
223 <211>	LENGTH:	242					
224 <212>	TYPE:	DNA					
225 <213>	ORGANISM:	Homosapiens					
227 <220>	FEATURE:						
228 <221>	NAME/KEY:	misc_feature					
229 <222>	LOCATION:	(0)...(0)					
230 <223>	OTHER INFORMATION:	N=A,T,C or G					
232 <400>	SEQUENCE:	11					

RAW SEQUENCE LISTING

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Input Set : A:\029001.txt

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W--> 233 actctcnngc ctctcaccga agatacgccg caaggactgg cgngaacann ggcgcgtgga 60
 W--> 234 ctatcnctaa agggtctccn acnacgtcca nccggacnag ctgacctcg ttcncnaag 120
 W--> 235 cgtgaaactg aaggccgtg aaaccntcnt gttcgctng atcacctact agtcgcgc 180
 W--> 236 cnngcgcgac aggatcaacg ccaaggtat ggccgatccc cgcctggcgt cgtcgatgga 240
 237 tc 242
 239 <210> SEQ ID NO: 12
 240 <211> LENGTH: 552
 241 <212> TYPE: DNA
 242 <213> ORGANISM: Homosapiens
 244 <400> SEQUENCE: 12
 245 gatccgctcg atgcccaggc ccagtacagc gaactgttcg cccatggccg egccacgtca 60
 246 ctgttgctat tcgaacatgt tcacggtaa tcccggtacc gggccaggc gatggggac 120
 247 ctgctggcgc agtacgagca gcacggtttg cagttaaaca gccgcgaatt accggaccac 180
 248 ctgcccgtgt atctggatg cctgtcgacg ctgcccgaag gcaagccgt ggaagggttt 240
 249 aaagatatacg cgccgattct ggcattgtct agcgcgcgtc tgcaacageg tgaaagccgt 300
 250 tatgccgtga tgtttgcgt gctgctgaaa ttggccgata ccgtatcgca cagcgacaaa 360
 251 gtggcggaaa aaattggca cgaagccgcg gatgatacgc cgccaggcgt ggtgctgtt 420
 252 tggaaagaag agcaggtaa attctttgtc gacaaggct gccgcattc agcaatcact 480
 253 gtcatcagc gtcgcttgc cggtgcgcgc gcccgcata atctgaatat cctcggtgag 540
 254 aggctggaga gt 552
 256 <210> SEQ ID NO: 13
 257 <211> LENGTH: 265
 258 <212> TYPE: DNA
 259 <213> ORGANISM: Homosapiens
 261 <220> FEATURE:
 262 <221> NAME/KEY: misc_feature
 263 <222> LOCATION: (0)...(0)
 264 <223> OTHER INFORMATION: N=A,T,C or G
 266 <400> SEQUENCE: 13
 W--> 267 gatcctnaca cantagcccg tggacgcatt tgcgtcgacc ctcatangga agcgatacga 60
 W--> 268 ggcgggttnaa agtgaacatc cggcggcac ggcagcgac cctccgtca ccgtcngcgc 120
 W--> 269 agtacttcct cgggtcgccg cgccttagcac tctgcgcgt gacatcaanc cgtgaaccca 180
 W--> 270 cgggagactt tgcggccgna agggatgagt ccactattag atgacgcgt gctacgagcc 240
 W--> 271 natcctcggt ganaagctgg agagt 265
 273 <210> SEQ ID NO: 14
 274 <211> LENGTH: 317
 275 <212> TYPE: DNA
 276 <213> ORGANISM: Homosapiens
 278 <220> FEATURE:
 279 <221> NAME/KEY: misc_feature
 280 <222> LOCATION: (0)...(0)
 281 <223> OTHER INFORMATION: N=A,T,C or G
 283 <400> SEQUENCE: 14
 W--> 284 gatccggccn cgcacganct taccggtnaa aacttccnccn ccnataatat ttgcccgcg 60
 W--> 285 agccgcctcg angctctcg cgtaaactccg gatgcacggg ggaccgtgac ggttgtantg 120
 W--> 286 ccctggcttt ttcagcnga aatctgcaca gcatcttc gatcgatctg gccgcagggtgg 180
 W--> 287 ggcggcncaa aacgggtggc atctccaaac cgcaggaaacg tgttttgcag gatgtcgaac 240
 W--> 288 atcatccacg cttcggttncc caacggctac ttccggccgtt accggggccat gtcatcctcg 300
 W--> 289 gtganaagct gganant 317

FYI:

Please Note:

Use of n and/or Xaa have been detected in the Sequence Listing. Please review the Sequence Listing to ensure that a corresponding explanation is presented in the <220> to <223> fields of each sequence which presents at least one n or Xaa.

VERIFICATION SUMMARY
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Input Set : A:\029001.txt
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L:186 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:188 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:189 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:190 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:191 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:192 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:193 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:194 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:195 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:196 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:9
L:213 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:10
L:216 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:10
L:217 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:10
L:218 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:10
L:219 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:10
L:233 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:11
L:234 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:11
L:235 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:11
L:236 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:11
L:267 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:13
L:268 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:13
L:269 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:13
L:270 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:13
L:271 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:13
L:284 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:14
L:285 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:14
L:286 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:14
L:287 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:14
L:288 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:14
L:289 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:14
L:303 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:15
L:304 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:15
L:305 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:15
L:306 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:15
L:307 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:15
L:321 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:16
L:322 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:16
L:323 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:16
L:340 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:17
L:340 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:17
L:340 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:17
L:342 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:17
L:342 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:17
M:340 Repeated in SeqNo=17
L:343 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:17
L:343 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:17
L:344 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:17
L:344 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:17

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L:345 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:17
L:345 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:17
L:346 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:17
L:346 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:17
L:363 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:18
L:363 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:18
L:363 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:18
L:364 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:18
L:364 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:18
M:340 Repeated in SeqNo=18
L:365 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:18
L:365 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:18
L:381 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:382 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:389 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:390 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:393 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:394 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:401 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:402 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:409 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:410 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:413 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:414 M:341 W: (46) "n" or "Xaa" used, for SEQ ID#:19
L:446 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:446 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:446 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:20
L:450 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:450 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
M:340 Repeated in SeqNo=20
L:452 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:452 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:454 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:454 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:456 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:456 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:460 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:460 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:462 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:462 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:464 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:464 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:466 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:466 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:468 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:468 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:470 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:20
L:470 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:20
L:552 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:22

VERIFICATION SUMMARY

PATENT APPLICATION: US/09/484,577A

DATE: 02/13/2001

TIME: 14:31:58

Input Set : A:\029001.txt

Output Set: N:\CRF3\02132001\I484577A.raw

L:552 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:22
L:552 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:22
L:556 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:22
L:556 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:22
M:340 Repeated in SeqNo=22
L:558 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:22
L:558 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:22
L:560 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:22
L:560 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:22
L:562 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:22
L:562 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:22
L:657 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:24
M:340 Repeated in SeqNo=24
L:697 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:25
M:340 Repeated in SeqNo=25
L:722 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:26
M:340 Repeated in SeqNo=26
L:762 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:27
M:340 Repeated in SeqNo=27
L:802 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:28
M:340 Repeated in SeqNo=28
L:923 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:30
M:340 Repeated in SeqNo=30
L:1034 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:32
M:340 Repeated in SeqNo=32
L:1145 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:34
M:340 Repeated in SeqNo=34
L:1175 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:35
M:340 Repeated in SeqNo=35
L:1201 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:36
M:340 Repeated in SeqNo=36
L:1243 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:37
M:340 Repeated in SeqNo=37
L:1285 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:38
M:340 Repeated in SeqNo=38
L:1357 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:40
M:340 Repeated in SeqNo=40
L:1408 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:42
M:340 Repeated in SeqNo=42
L:1459 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:44
M:340 Repeated in SeqNo=44
L:1481 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:45
M:340 Repeated in SeqNo=45
L:1501 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:46
M:340 Repeated in SeqNo=46
L:1523 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:47
M:340 Repeated in SeqNo=47
L:1545 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:48
M:340 Repeated in SeqNo=48

VERIFICATION SUMMARY

PATENT APPLICATION: US/09/484,577A

DATE: 02/13/2001

TIME: 14:31:58

Input Set : A:\029001.txt

Output Set: N:\CRF3\02132001\I484577A.raw

L:2021 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:60
M:340 Repeated in SeqNo=60

L:2084 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:62
M:340 Repeated in SeqNo=62

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Robinson
H.

RAW SEQUENCE LISTING
PATENT APPLICATION: US/09/319,860A

DATE: 02/13/2001
TIME: 14:30:41

Input Set : A:\09-31981.app
Output Set: N:\CRF3\02132001\I319860A.raw

ENTERED

3 <110> APPLICANT: Grant, Peter J
 5 <120> TITLE OF INVENTION: Atherothrombotic Disorders: Mutation in the Factor XIII
 6 Gene
 8 <130> FILE REFERENCE: Sequence Listing SN 09/319,860
 10 <140> CURRENT APPLICATION NUMBER: 09/319,860A
 11 <141> CURRENT FILING DATE: 1999-06-11
 13 <150> PRIOR APPLICATION NUMBER: PCT/GB97/03435
 14 <151> PRIOR FILING DATE: 1997-12-12
 16 <150> PRIOR APPLICATION NUMBER: GB 9625934.6
 17 <151> PRIOR FILING DATE: 1996-12-13
 19 <160> NUMBER OF SEQ ID NOS: 3
 21 <170> SOFTWARE: PatentIn Ver. 2.1
 23 <210> SEQ ID NO: 1
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 25 <212> TYPE: DNA
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 29 <223> OTHER INFORMATION: Description of Artificial Sequence: PCR primer
 30 sequence
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 33 acccagagtgtggggaa 19
 36 <210> SEQ ID NO: 2
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 39 <213> ORGANISM: Artificial Sequence
 41 <220> FEATURE:
 42 <223> OTHER INFORMATION: Description of Artificial Sequence: PCR primer
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 46 gaccttgtaa agtcaaaaat gtc 23
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 51 <212> TYPE: DNA
 52 <213> ORGANISM: Homo sapiens
 54 <220> FEATURE:
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 56 <222> LOCATION: (1)..(42)
 57 <223> OTHER INFORMATION: exon 2 of Factor XIIIa
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VERIFICATION SUMMARY

PATENT APPLICATION: US/09/319,860A

DATE: 02/13/2001

TIME: 14:30:42

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Output Set: N:\CRF3\02132001\I319860A.raw